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INTRODUCTION

While an increasing number of women at risk for ovarian cancer are being identified through awareness efforts and risk assessment programs, a gap still exists in the known psychological and physical sequelae of preventive surgery options offered to these women. To meet the needs of women seeking information about the effects of prophylactic oophorectomy and for those who will undergo the procedure, this study will provide significant information on the broader quality of life domains and physiologic changes following surgery. In order to make informed decisions about their choices, women considering prophylactic oophorectomy need scientific data on the hormonal and other physiologic consequences of surgery, and on the potential alterations in their emotional and social well being. They also need the opportunity to choose from an array of coping strategies to manage their health decisions. Ovarian cancer advocacy groups have voiced their support for research that advances not only ovarian cancer prevention and treatment but also quality of life. Studying multidimensional quality of life issues will contribute to the knowledge base about the short and long-term effects on physical, emotional, cognitive, sexual and social functioning following oophorectomy and will contribute to the development of optimum medical and alternative therapy strategies to deal with post-surgical sequelae. As important, it will also identify issues and needs faced by women who make the choice not to undergo surgery.

BODY

The following describes the progress during the past year associated with each task in the Statement of Work.

Task 1: Creation of Participant Advisory Board

We continue with an informal advisory approach, holding consultations by telephone or on a person-to-person basis. Collaborative efforts have been initiated with new gynecologic oncology staff, Dr. Norman Rosenblum and Dr. Mitchell Edelson at Fox Chase Cancer Center, to identify women contemplating prophylactic surgery. Dr. Marcelle Shapiro, physician and participant in the Family Risk Assessment Program, has offered her services in an advisory capacity. We continue our networking relationship to several local advocacy groups, including the Philadelphia chapter of the National Ovarian Cancer Coalition and The Sandy Rollman Ovarian Cancer Foundation, Inc.

Task 2: Selection of Survey Instruments

There have been no changes in the survey instruments previously approved by the IRB. Outcome variables include physical functioning, menopausal symptoms, body image, sexual functioning, anxiety, depression, and use of pharmaceutical, dietary and alternative therapies. The instruments being used are as follows:

1. The NSABP BCPT Quality of Life Questionnaire. This instrument was used by over 13,000 women in the Tamoxifen prevention trial. It includes the Medical Outcomes Study (MOS) 36-item short form, a generic measure of health-related QOL, the Center for Epidemiologic Studies-Depression Scale, used widely in

- community epidemiologic studies, the MOS sexual problems scale, and a 43-item symptom checklist of commonly reported physical and psychologic symptoms, as well as symptoms associated with the menopause, including the domains of vasomotor symptoms, vaginal dryness, sexual functioning, sleep disturbance and cognitive functioning. Sleep patterns and sleep quality may be disrupted by surgical menopause. This questionnaire is collected at all time points.
2. Post-Surgical Expectations Questionnaire. The NSABP BCPT Quality of Life Questionnaire has been modified to assess women's expectations of menopausal symptoms they anticipate experiencing following oophorectomy. It includes an open-ended response format as well as a Likert-type summary scale of symptoms. This questionnaire is only assessed at baseline, prior to surgery.
 3. Fallowfield Sexual Activity Questionnaire (SAQ). This tool is a validated measure for describing the sexual functioning of women in terms of activity, pleasure and discomfort. It was developed to investigate the impact of long-term Tamoxifen usage on the sexual functioning of women at high risk of developing breast cancer. This measure is collected at all time points.
 4. Self Concept Scale. This 10-item scale assesses the participants' satisfaction with different areas of their body and their overall weight. Persons undergoing oophorectomy may experience an alteration in their perception of their body image, which may affect their psychosocial status and intimate relationships. This scale was developed by Dr. David Cella, (Director, Center on Outcomes Research and Education, Evanston Hospital) through his work with breast cancer patients. It is collected at all time points.
 5. Medical/Dietary Supplement Survey. This survey elicits use of hormone replacement therapy, dietary supplements, micronutrients, as well as exercise, yoga, meditation, and other forms of coping strategies. The survey has been piloted among 48 women in the FRAP program for feasibility and ease of administration. Overall, we found that 89% of the women surveyed took some form of dietary supplement. It is collected at all time points.
 6. Post-Surgery Satisfaction Questionnaire. Patients' levels of satisfaction with oophorectomy will be assessed using three items rated on a 5-point Likert-type scale. Scores from the three items will be combined to form a composite index of satisfaction. It is collected at all post-surgery time points.
 7. Medical Outcomes Survey. This survey will capture information on new medical diagnoses, procedures, and screening exams at the 12-month follow-up time point. It is adapted from our current FRAP annual follow-up questionnaire.

Task 3: Development of a Recruitment Strategy

Enrollment into the surgery arm has been steady with a total of 30 women recruited. There are 11 women in the control group and one additional woman is noncompliant. We continue to identify potential candidates through the Family Risk Assessment Program (FRAP) at cancer risk counseling and clinical exam appointments. It is our experience that the decision making process can be quite lengthy for many women who consider prophylactic surgery, which has challenged us in identifying a point in time when the control arm candidates decide against surgery. A recruitment letter to *BRCA1* and *BRCA2* mutation carriers that are screened in our

FRAP clinic, (and have not had prophylactic surgery), has been submitted to our IRB. After approval we will telephone these women to explain the research study, in an effort to bolster control arm enrollment.

Task 4: Creation of Data Entry Screens, Data Editing Program

This task is completed and data entry is a smoothly flowing process. Data is entered promptly and close consultation between the project manager and data entry clerk has served to oversee data editing review. A series of edit checks and quality assurance measures take place on a routine basis whenever data is entered into our bioinformatics system.

Task 5 & 6: Conduct Baseline & Follow-up Surveys

Surveys are sent in packets with an instruction cover letter and postage paid return envelope. The project manager receives an email notice to trigger the sending of packets. Reminder postcards and personal phone calls are made when warranted to alert participants of overdue surveys.

Task 7: Data entry, data analysis

Data entry is up to date. The project manager tracks study accrual and questionnaire completion on a biweekly basis. Interim short-term analysis of one-month follow-up data on 22 women in the surgery arm was performed by Dr. Carolyn Fang for a poster presentation at the American Society of Preventive Oncology meeting in March 2002. Dr. Fang is an investigator in the behavioral research center at Fox Chase Cancer Center.

Task 8: Report, manuscript preparation

The required application for ongoing review by the Research Review and IRB committees at Fox Chase Cancer Center was prepared and approved on 6/14/02. Insufficient data exists at this time for manuscript preparation.

KEY RESEARCH ACCOMPLISHMENTS

- Dr. Carolyn Fang's interim short-term analysis comparing baseline (pre-surgery) assessments of 22 women with their one-month post-surgery assessments showed short-term decrements in satisfaction with, and frequency of, sexual activity. There were no significant changes in psychological distress, self-concept, or the *total* number of physical symptoms observed after surgery. However, a higher percentage of women reported certain physical symptoms, such as fatigue, decreased appetite, and night sweats, after surgery than before it, and those women were more distressed.

REPORTABLE OUTCOMES

A poster entitled, "Psychosocial and Sexual Functioning following Prophylactic Oophorectomy," was prepared by Dr. Carolyn Fang and presented at the annual meeting of the American Society of Preventive Oncology (ASPO), March 2002, in Bethesda, MD.

CONCLUSIONS

The short-term interim analysis performed by Dr. Carolyn Fang lays groundwork for developing interventions to prepare at-risk women for issues arising from prophylactic surgery, but sufficient numbers of study participants must be accrued for meaningful outcomes. As more is learned about the nature of physical and psychological symptoms following surgery we will be better equipped to educate and counsel at-risk women regarding the quality of life issues related to preventive clinical options.

REFERENCES

None

APPENDICES

None